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CLAIMS

1. A compound according to the general formula:

$$R_5$$
— $(CH_2)_n$ — O — R_4
 R_5 — CH_2 - R_1

or a pharmaceutically acceptable salt thereof, wherein:

R₁ is independently selected from: carboxylic acid (-CO₂H); phosphonic acid (-PO(OH)₂); phosphamic acid (-PO(OH)NH₂); sulphonic acid (-SO₂OH); hydroxamic acid (-CONHOH); oxamic acid (-NHCOCO₂H); and malonamic acid (-NHCOCH₂CO₂H), or any other possible bioisosteric equivalent of the groups above;

R₂ and R₃ are the same or different and independently selected from: chlorine; bromine; iodide; C₁₋₄ alkyl, said alkyl, or a bioisosteric equivalent optionally substituted with 0, 1, 2 or 3 groups of R^a which groups may be the same or different;

 R_4 and R_6 are the same or different and independently selected from: hydrogen; halogen; C_{1-4} alkyl; or a bioisosteric equivalent optionally substituted with 0, 1, 2 or 3 groups of R^a which groups may be the same or different;

R₅ is selected from: C₆₋₁₀ aryl; C₁₋₉ heteroaryl, said aryl; and heteroaryl optionally substituted with 0, 1, 2, or 3 groups of R^b which groups may be the same or different;

Ra represents fluorine or chlorine;

R^b represents a member selected from the group of: halogen; -CN; -CO₂H; -CHO; -NH₂; C₁₋₄ alkyl; C₂₋₄ alkenyl; C₂₋₄ alkynyl; C₁₋₄ alkoxy; C₂₋₄ alkenoxy; C₂₋₄ alkynoxy; C₁₋₄ alkylthio; C₂₋₄ alkenylthio; C₂₋₄ alkynylthio; C₆ aryl; C₁₋₅ heteroaryl; C₃₋₆ cycloalkyl; -NH(C₁₋₄); -N(C₁₋₄)₂; -NH(C₆ aryl); -N(C₆ aryl)₂; -NH(C₁₋₅ heteroaryl)₂ or a bioisosteric equivalent;

n is an integer of 1, 2 or 3;

included for the variables above are all the possible stereoisomers thereof; prodrug ester forms thereof; and radioactive forms thereof.

- 2. A compound according to claim 1 wherein R₁ is carboxylic acid (-CO₂H).
- 3. A compound according to claim 1 or 2 wherein R₂ and R₃ is bromine or chlorine.
- 4. A compound according to any one of claims 1 to 3 wherein R₄ is isopropyl and R₆ is hydrogen.
- 5. A compound according to any one of claims 1, 2 or 4 wherein R₂ and R₃ is bromine.
- 6. A compound according to any one of claims 1 to 5 which is:
 - {4,6-Dibromo-5-[3-isopropyl-4-(naphthalen-2-yl-methoxy)phenoxy]indan-1-yl}-acetic acid;
 - {4,6-Dibromo-5-[4-(4-fluorobenzyloxy)-3-isopropylphenoxy]indan-1-yl}acetic acid;
 - {4,6-Dibromo-5-[3-isopropyl-4-(5-methylisoxazol-3-ylmethoxy)phenoxy]indan-1-yl}acetic acid;
 - {4,6-Dibromo-5-[3-isopropyl-4-(pyridin-2-yl-methoxy)phenoxy]indan-1-yl}acetic acid;
 - 4,6-Dibromo-5-[3-isopropyl-4-(5-phenyl-[1,2,4]oxadiazol-3-ylmethoxy)phenoxy]-indan-1-yl}acetic acid;
 - 4-[4-(4,6-Dibromo-1-carboxymethyl-indan-5-yloxy)-2-isopropylphenoxymethyl]-benzoic acid;
 - (4,6-Dibromo-5-4-[2-(1*H*-indol-2-yl)ethoxy]-3-isopropylphenoxy}indan-1-yl) acetic acid;
 - (4,6-Dibromo-5-[3-isopropyl-4-(5-thiophen-3-yl-[1,2,4]oxadiazol-3- yl-methoxy)-phenoxy[indan-1-yl] acetic acid;
 - {5-[4-(4-Amino-6-phenylamino[1,3,5]triazin-2-ylmethoxy)-3-isopropylphenoxy]-4,6-dibromoindan-1-yl}acetic acid;

- {4,6-Dibromo-5-[3-isopropyl-4-(5-methyl-2-phenyloxazol-4-ylmethoxy)phenoxy]-indan-1-yl}acetic acid;
- {4,6-Dibromo-5-[4-(3,5-dimethylisoxazol-4-ylmethoxy)-3-isopropylphenoxy]-indan-1-yl}acetic acid;
- and pharmaceutically acceptable salts thereof, and stereoisomers thereof.
- 7. A compound according to any one of claims 1 to 6, which have one or more asymmetric centers and can exist in the form of racemates, single and multiple enantiomers, as individual diastereomers, with all possible isomers, and mixtures thereof.
- 8. A compound according to any one of claims 1 to 7 for use in medical therapy.
- 9. A pharmaceutical composition comprising an effective amount of a compound according to any one of claims 1 to 7 or a pharmaceutically effective salt thereof, together with a pharmaceutically acceptable carrier.
- 10. A method for preventing, inhibiting or treating a disease which is dependent on the expression of a T₃ regulated gene or associated with metabolic dysfunction, which comprises administering to a patient in need of treatment a therapeutically effective amount of a compound according to any one of claims 1 to 7.
- 11. The method according to claim 10 wherein the disease is selected from cardiac arrhythmias, thyrotoxicosis, subclinical hyperthyrodis, certain skin disorders, and certain liver diseases.
- 12. The method according to claim 11 wherein the disease is a skin disorder or skin disease.
- 13. The method according to claim 12 wherein the skin disorder or skin disease is selected from: keloids, lichen planus, ichtyosis, acne, psoriasis, Dernier's disease, eczema, atopic dermatitis, chloracne, pityriasis, and hirsuitism.
- 14. The method according to claim 11 wherein the disease is a liver disorder or liver disease.

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- 15. The method according to claim 14 wherein the liver disorder or liver disease is selected from: chronic alcoholism, acute hepatitis, chronic hepatitis, hepatitis C-induced liver cirrhosis, and liver fibrosis.
- 16. A method to treat certain skin disorders or diseases by the use of a compound according to any one of claims 1 to 7 in a combination with a retoid or a Vitamin D analog.
- 17. The use of a compound according to any one of claims 1 to 7 in the preparation of a medicament for the treatment of a disease or disorder which is dependent on the expression of a T₃ regulated gene.
- 18. The use according to claim 17 wherein the disease or disorder is cardiovascular disorder, thyrotoxicosis, subclinical hyperthyrodism, certain skin disorders and certain liver diseases.
- 19. The use according to claim 17 wherein the disease or disorder is selected from atrial fibrillation, ventricular tachycardia and ventricular fibrillation.
- 20. The use according to claim 17 wherein the disease or disorder is selected from thyrotoxicosis, subclinical hyperthyrodism and other related endocrine disorders, related to thyroid hormone.
- 21. The use according to claim 17 wherein the disease is a liver disorder or liver disease.
- 22. The use according to claim 21 wherein the liver disorder or liver disease is selected from: chronic alcoholism, acute hepatitis, chronic hepatitis, hepatitis C-induced liver cirrhosis, and liver fibrosis.
- 23. The use of a compound according to any one of claims 1 to 7 in the preparation of a medicament for the treatment of anoxic tissue damage.
- 24. The use of a labeled compound, according to any one of claims 1 to 7, as a diagnostic agent.